

HAIR TREATMENT AGENT, SHAMPOO AND HAIR COLORING AGENT

FIELD OF THE INVENTION

Present invention relates to a hair treatment agent
5 including a hair rinse, hair conditioner, hair treatment,
hair treatment cream, PPT(polypeptide) treatment, hair
cream, hair spray, hair wax and the like, a shampoo and
a hair coloring agent. More specifically, the present
invention relates to
10 a hair treatment agent imparting excellent gloss and
springness to hairs, making hairs smooth and improving
combability of hairs, preventing formation of split hairs,
and improving manageability of hairs, particularly a hair
treatment agent exhibiting an excellent effect of
15 preventing discoloration of dyed hairs, in addition to the
above mentioned effects, when used for dyed hairs;
a shampoo imparting gloss to hairs, making hairs smooth
and improving combability of hairs and improving
manageability of hairs for the shampooed hair; and
20 a hair coloring agent belonging to oxidation type hair dyes,
capable of dyeing hairs uniformly, giving less damage to
hairs on dyeing, imparting gloss, springness and improved
combability to dyed hairs, and having excellent shampoo
fastness.

25

BACKGROUND OF THE INVENTION

Conventionally used hair treatment agents such as hair
rinses, hair conditioners, hair treatments and the like

contain, as the conditioning agents, cationic surfactants
silicones, oily materials, hydrolyzed proteins and the
like.

However, cationic surfactants soften hairs too much
5 and deteriorate springness of hairs, although cationic
surfactants exhibit good sorption action. Hydrophobic
materials such as silicones and oily materials used in
cosmetics are easily sorbed by little damaged hairs, that
is, highly hydrophobic hairs, but hardly sorbed by damaged
10 hairs having high hydrophilicity given by the hydrophilic
groups come out to the surface of hairs caused by the damage.
Consequently, the silicones and oily materials cannot exert
sufficiently their properties to the damaged hairs.

Hydrolyzed proteins exhibit good sorption action, and
15 particularly have excellent effects
of recovering deteriorated moisture-keeping actions of
damaged hairs, the deterioration being caused by elution
of protein components of hairs by chemical treatments such
as permanent waving, dyeing or the like or by daily shampoos,
20 and
of imparting moisture feeling to hairs. However,
hydrolyzed proteins have problems that they compare
unfavorably in imparting springness to hairs with high
molecule silicones, and they tend to give stickiness when
25 contained in high concentration.

Shampoos conventionally used contain as their major
ingredients anionic surfactants such as alkyl sulfate,

polyoxyethylenealkyl sulfate and the like, nonionic surfactants such as polyoxyethylenealkylether, fatty acid alkylol amide and the like, and ampholytic surfactants such as alkylbetaine, alkylamineoxide and the like, in alone
5 or in the mixture thereof. Furthermore, a shampoo is proposed which contains cationic surfactants to give soft and moist feeling to hairs.

However, if washing hair with the shampoo containing above described surfactants, sebum and other fat existing
10 on the surface of hairs are excessively washed away. Consequently, the gloss and moisture of hairs are lost, the feeling of hairs is extremely deteriorated like overly dried or roughly dried hairs, and split hair and fractured hair are formed due to deterioration of combability and
15 brushing ability.

Therefore, even in the above described shampoos, as being proposed other cosmetics for hair, an improvement of blending a silicone compound called as silicone oil and a polypeptide is proposed to exert their excellent
20 properties together. Because the silicone oil is excellent in its extensibility, imparting gloss and luster to hair and hair protecting action resulting from imparting water-repelling property to hair, and because the polypeptide is excellent in substantivity to hair and
25 protective and moisture-keeping actions based on film formation.

However, because silicone oils are hydrophobic (lipophilic) compounds and polypeptides are hydrophilic

compounds, they are poor in compatibility with each other. Accordingly, when they are used in combination, commodity value as shampoo is apt to be impaired because the emulsion stability is low and separation is easy to occur. Also, because the polypeptides are hard to attach to a portion of the hair which has contacted the silicone oils in advance of the polypeptides, and conversely because the silicone oils are hard to attach to a portion of hair which has contacted the polypeptides in advance of the silicone oils, the combination use of the silicone oils and the polypeptides has a problem that the properties of them cannot be sufficiently exerted.

Hair coloring agents include temporally hair coloring agents, semi-permanent hair coloring agents, permanent hair coloring agents and the like, and have been widely used for beauty culture.

As the permanent hair coloring agents, an oxidation type hair coloring agent consisting of a first agent containing an oxidation dye (a dye intermediate) and a second agent containing an oxidant such as hydrogen peroxide are widely used, since various color tones are obtained and the coloring power is excellent.

However, the oxidation type hair coloring agent has problems that it takes longer time for the color reaction on hair than acidic hair coloring agents, temporally hair coloring agents containing an acid dye as the main ingredient, since the first agent containing an oxidation dye and the

second agent containing an oxidant need to be mixed before applying the agent on hair, and that it hardly gives uniform dyeing.

In addition, the oxidation type hair coloring agent usually contains an alkali in order to facilitate uniform penetration of an oxidation dye such that the pH is as high as 10 or more, and accordingly it has the defect that it is stimulative and the hair is easy to be damaged and the protein component in the hair is easy to be eluted. As the results, there are problems that the dyed hair is overly dried, combability is deteriorated, discoloration becomes large caused by daily shampoo which elutes dye in the damaged hair, and the like.

In order to solve the problems and to improve uniform dyeability of the hair coloring agent, it is proposed to contain hydrolyzed proteins (peptide), derivatives thereof, silicones and the like in the hair coloring agent (for example, JP-A 61-55887, JP-B 3-63528, JP-B 4-4289, JP-A 5-271040).

However, the above-mentioned hydrolyzed proteins and derivatives thereof are easily washed away by shampoo after hair dyeing, and do not impart sufficiently satisfied shampoo fastness to the dyed hair, although they are sorbed to the dyed hair and impart the hair moisture feeling and gloss. Silicone is difficult to be sorbed by damaged hair in which hydrophilic groups come out to the surface, although it is easily sorbed by undamaged hair. Further, silicone has a problem that dyeability of the hair coloring agent

is lowered when the hair coloring agent contains large amount of silicone.

To solve above described problems regarding hair treatment agent, shampoo and hair coloring agent, inventors of the present invention have developed a silylated peptide which has silyl group added to hydrolyzed peptide by addition reaction. And they also have tried to use it for a hair treatment agent, a shampoo and a hair coloring agent to co-exert both of the properties of hydrolyzed peptide such as imparting gloss, moisture feeling and manageability to hairs, and of the properties of silyl group such as imparting gloss to hairs and making hairs smooth (for example, JP-A 8-81338, JP-A 2002-302648, JP-A 8-81339, JP-A 8-157344).

However, because the number of silyl group is only one which is added to the peptide portion of the above described silylated peptide, there is a problem that the property imparting smoothness on the surface of hairs is inferior to high molecule silicone.

Furthermore, regarding shampoo, because the peptide portion having hydrophilicity is large, there is another problem that the amount sorbed on hairs becomes less due to being washed away in shampooing. Regarding hair coloring agent, shampoo fastness of dyed hair is not satisfactory good. In these reasons, the above described silylated peptide is not yet sufficiently satisfied.

Therefore, the object of the present invention is to

provide a hair treatment agent in which the above-mentioned problems of prior art are solved, which imparts good gloss and springness to hairs, improves combability of hairs, prevents formation of split hairs, improves manageability
5 of hairs, and particularly, is excellent in a property which prevents discoloration of hairs due to shampoo, when used for dyed hairs, in addition to the properties mentioned above.

Another object of the present invention is to provide
10 a shampoo which fully co-exerts the excellent properties of silicone compound and peptide, and which imparts good gloss to hairs, makes hairs smooth and improves combability of hairs, and improves manageability of hairs.

Still another object of the present invention is to
15 provide a hair coloring agent which dyes hairs uniformly, gives little damage to hairs on dyeing, and imparts good gloss, springness, combability and the like to hairs, and makes shampoo fastness of dyed hairs excellent.

20 The present inventors have been intensively studied to solve the above-described problems, and consequently have found that a hair treatment agent, a shampoo and a hair coloring agent comprising a specific silylated peptide-silane compound copolymer composition can attain
25 the objects.

That is, they have found that a hair treatment agent prepared by comprising the specific silylated peptide-silane compound copolymer composition can attain

the above object to provide properties such as imparting good gloss and springness to hairs, improving combability of hairs, preventing formation of split hairs, improving manageability of hairs, and preventing discoloration of hairs due to shampoo, when used for dyed hairs.

They have also found that a shampoo prepared by comprising the specific silylated peptide-silane compound copolymer composition can attain the object to provide properties such as imparting good gloss to hairs, making hairs smooth and improving combability of hairs and improving manageability of hairs.

They have further found that a hair coloring agent prepared by comprising the specific silylated peptide-silane compound copolymer composition can attain the object to provide properties such as dyeing hairs uniformly, giving little damage to hairs on dyeing, and imparting good gloss, springness, combability and the like to hairs, and making shampoo fastness of dyed hairs excellent.

Base on these founded facts, the present invention was completed.

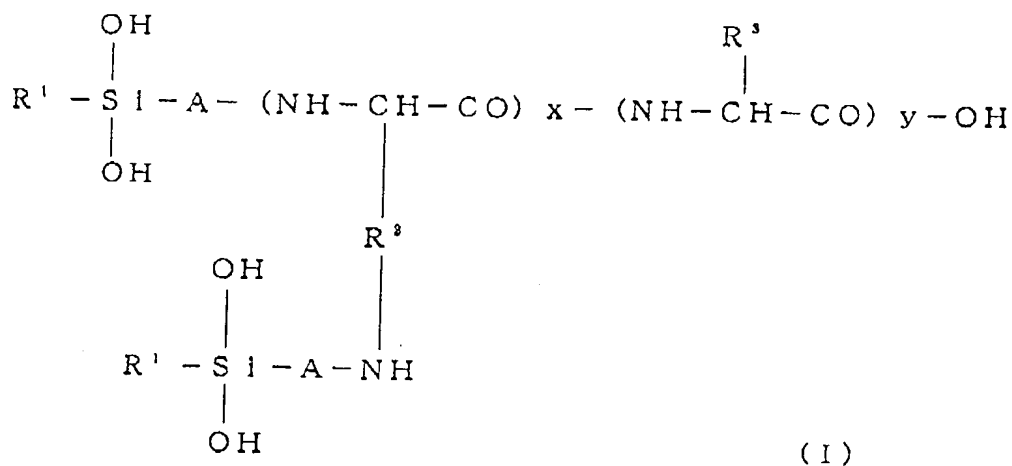
SUMMARY OF THE INVENTION

The present invention provides a hair treatment agent, a shampoo or a hair coloring agent comprising a silylated peptide-silane compound copolymer composition which is produced by the following process and has viscosity in a range from 500 to 20000 mPa·s in 70% of solid content

concentration of said copolymer composition at 20 °C,
 wherein said process comprises:

polycondensing one or more kind selected from silylated
 peptides represented by the general formula (I) with one
 5 or more kind selected from silane compounds represented
 by the general formula (II) in an aqueous solution in a
 range of reaction molar ratio of said silylated peptide
 to said silane compound from 1:1 to 1:100 to form a
 polycondensed polymer, and then
 10 adding by addition reaction a silane compound represented
 by the general formula (III) in an aqueous solution to said
 polycondensed polymer;

wherein said silylated peptide represented by the following
 general formula (I) is

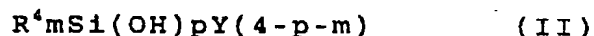


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 20 , in which R¹ represents a hydroxy group or an alkyl group
 having 1 to 3 carbon atoms, R² represents a residual group
 of a side chain obtained by removing the terminal end amino
 group of a basic amino acid having an amino group at the

end of a side chain, R^3 represents a side chain of an amino acid other than R^2 , A is a connecting moiety and represents at least one group selected from the group consisting of $-CH_2-$, $-(CH_2)_3-$, $-(CH_2)_3OCH_2CH(OH)CH_2-$, $-(CH_2)_3S-$,

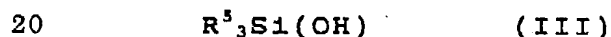
5 $-(CH_2)_3NH-$ and $-(CH_2)_3OCOCH_2CH_2-$, and x is from 0 to 50, y is from 1 to 100 and $x+y$ is from 1 to 100, wherein x and y represent only the number of amino acid units and do not represent the order of amino acid sequence;

wherein said silane compound represented by the following
10 general formula (II) is



, in which m represents an integer from 0 to 2, p represents an integer from 2 to 4, $m+p$ is not more than 4, R^4 represents an organic group in which a carbon atom is directly connected
15 to the silicon atom and R^4 's of m may be the same or different, and Ys of $(4-p-m)$ represent an alkoxy group or hydrogen atom; and

wherein silane compound represented by the following general formula (III) is



, wherein three R^5 's represent organic groups in which a carbon atom is directly connected to the silicon atom and three R^5 's may be the same or different.

The letters of x and y used in the above described
25 general formula (I) and of m, p and $(4-p-m)$ used in the above described general formula (II) are the substitute characters.

DETAILED DESCRIPTION OF THE INVENTION

The silylated peptide-silane compound copolymer composition used for the hair treatment agent, shampoo or hair coloring agent of the present invention may be produced
5 by the methods, for example, disclosed in JP-A 2001-48732 and in JP-A 2001-48775. A method for producing said silylated peptide-silane compound copolymer composition is described in more detail hereinafter.

10 A silylated peptide represented above described general formula (I), which is one of components of the silylated peptide-silane compound copolymer composition, is easily produced in an aqueous solution by the methods disclosed in JP-A 8-59424 and JP-A 8-67608.

15 In the silylated peptide represented above described general formula (I), R^2 is a residual group of a side chain obtained by removing the terminal end amino group of a basic amino acid having an amino group at the end of a side chain. Examples of above described basic amino acid having an amino
20 group at the end of a side chain include lysine, arginine, hydroxylysine and the like. R^3 represents an amino acid side chain other than R^2 . Examples of the amino acid include glutamic acid, aspartic acid, alanine, serine, threonine, valine, methionine, leucine, isoleucine, tyrosine,
25 phenylalanine, proline, hydroxyproline and the like.

In the silylated peptide shown by the above described formula (I), x is from 0 to 50, preferably from more than 0 to not more than 10, y is from 1 to 100, preferably from

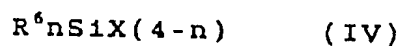
1 to 50, and $x+y$ is from 1 to 100, preferably from 2 to 50. If x is more than the above range, because the number of the silyl group bonding to the amino acid in the side chains increases, the sorption action to the hair that the peptide originally possesses is decreased. If y is more than the above range, because the proportion of the silyl group portion to the peptide portion is small, the effects based on the silyl group portion cannot be sufficiently exerted. If the $x+y$ is more than the above range, the sorbing ability and permeability to be exerted by the peptide is decreased in comparison with a low molecular weight peptide. The above described x , y and $x+y$ are theoretically an integer. However, in case that the peptide portion is derived from a hydrolyzed peptide, since the hydrolyzed peptide is obtained in the form of a mixture of those having a different molecular weight, the measured values become average values.

The peptide portion of the silylated peptide represented by the above described formula (I) is illustrated with natural peptide, synthetic peptide, a hydrolyzed peptide obtained by partially hydrolyzing a protein with an acid, an alkali or an enzyme, or the mixture thereof. Among of them, hydrolyzed peptide is preferably used according to the reason that the protein can be conveniently obtained and the number average molecular weight of the peptide portion can be easily controlled.

Examples of the hydrolyzed peptide include animal- or vegetable-derived protein such as collagen (including

gelatin of modified product thereof), keratin, silk fibroin (silk), sericin, casein, conchiolin, elastin, yolk protein and albumen protein of eggs such as fowl and duck, soybean protein, wheat protein, corn protein, rice (rice bran) protein and potato protein; yeast proteins separated from yeasts such as yeasts belonging to genera *Saccharomyces*, *Candida* and *Endomycopsis*, or yeasts such as so-called beer yeasts and sake yeasts; and peptides obtained by partially hydrolyzing microorganism-derived proteins such as proteins separated from fungi (*Basidiomycetes*) or *Chlorella* with acid, alkali or enzyme, or the mixture thereof; and the like.

A silane compound, which is another of components of the silylated peptide-silane compound copolymer composition used for the hair treatment agent, shampoo or hair coloring agent of the present invention, is represented by the above described general formula (II), and said compound is obtained by hydrolyzing a silane compound represented by the following general formula (IV) in aqueous solution;



, wherein n represents an integer from 0 to 2, R^6 represents an organic group in which a carbon atom is directly connected to the silicon atom and R^6 's of n may be the same or different, and X s of $(4-n)$ represent at least a group selected from the group consisting of a hydroxyl group, an alkoxy group and a halogen group. The letters of n and $(4-n)$ used in the above described general formula (IV) are the substitute

characters.

Specific examples of such a silane compound represented by the general formula (IV) include tetramethoxysilane, methyltrimethoxysilane, 5 methyldimethoxysilane, dimethyldimethoxysilane, phenyltrimethoxysilane, diphenyldimethoxysilane, hexyltrimethoxysilane, decyltrimethoxysilane, vinyltrimethoxysilane, 3-methacryloxypropyltrimethoxysilane, 10 3-methacryloxypropylmethyldimethoxysilane, N-(2-aminoethyl)-3-aminopropyltrimethoxysilane, N-(2-aminoethyl)-3-aminopropylmethyldimethoxysilane, 3-aminopropyltrimethoxysilane, N-phenyl-3-aminopropyltrimethoxysilane, 15 3-chloropropyltrimethoxysilane, 3-chloropropylmethyldimethoxysilane, 3-mercaptopropyltrimethoxysilane, 3-glycidoxypropyltrimethoxysilane, 3-glycidoxypropylmethyldimethoxysilane, 20 dimethyloctadecyl[3-(trimethoxysilyl)propyl] ammonium chloride, 3-(trimethoxysilyl)propylpolyoxyethylene (10) ether, tetraethoxysilane, methyltriethoxysilane, methyldiethoxysilane, dimethyldiethoxysilane, phenyltriethoxysilane, diphenyldiethoxysilane, 25 vinyltriethoxysilane, 3-methacryloxypropyltriethoxysilane, 3-methacryloxypropylmethyldiethoxysilane, N-(2-aminoethyl)-3-aminopropyltriethoxysilane,

N-(2-aminoethyl)-3-aminopropylmethyldiethoxysilane,
3-aminopropyltriethoxysilane,
3-chloropropyltriethoxysilane,
3-chloropropylmethyldiethoxysilane,
5 3-glycidoxypropyltriethoxysilane,
3-glycidoxypropylmethyldiethoxysilane,
3-isocyanatepropyltriethoxysilane, methyldichlorosilane,
methyltrichlorosilane, dimethyldichlorosilane,
phenyltrichlorosilane, diphenyldichlorosilane,
10 vinyltrichlorosilane,
3-chloropropylmethyldichlorosilane and the like.

The reaction of the silylated peptides represented
by the above described general formula (I) with the silane
compound represented by the general formula (II) is carried
15 out according to, for example, the following procedure:
Preparing an aqueous solution of the silylated peptides
represented by the above described general formula (I) to
become acidic with hydrochloric acid, sulfuric acid or the
like, or to become basic with an aqueous sodium hydroxide
20 solution, an aqueous potassium hydroxide solution or the
like;

Adding dropwisely the silane compound represented by the
general formula (IV) to said solution to form a silane
compound represented by the general formula (II) having
25 at least two hydroxyl groups directly connected to a silicon
atom by hydrolyzing an alkoxy group, a halogen group and
the like of the silane compound represented by the above
described general formula (IV);

Neutralizing the solution obtained thereby to polycondense a hydroxyl group of the organic silane compound having a hydrophilic group represented by the general formula (I) with a hydroxyl group of the silane compound represented
5 by the general formula (II).

Thus, a silylated peptide-silane compound copolymer composition is obtained.

The hydrolysis reaction generally proceeds in a state of pH 1 to 3. However, because certain kinds of silylated
10 peptide represented by the general formula (I) is likely to form insoluble materials in acidic state, when such silylated peptide is used, the hydrolysis is preferably conducted in the state of pH 10 to 11. When an alkoxysilane compound is used as the silane compound represented by the
15 general formula (IV), pH control is required only before adding dropwisely the alkoxysilane compound. However, when the reaction is conducted in basic state by using a halogenated silane compound or carboxysilane compound as the silane compound represented by the general formula (IV),
20 it is necessary to keep pH between 10 and 11 by adding an aqueous sodium hydroxide solution, an aqueous potassium hydroxide solution or the like due to pH of the reaction solution decreasing during the reaction. Further, when the reaction is conducted in acidic state by using an amino
25 silane compound as the silane compound represented by the general formula (IV), it is necessary to keep pH between 1 and 3 by adding diluted hydrochloric acid, diluted sulfuric acid or the like due to pH of the reaction solution increasing

during the reaction.

The reaction temperature is preferably from 30 to 60 °C, because the reaction does not proceed smoothly when the reaction temperature is too low, and because an alkoxy group or a halogen group of the silane compound represented by the above described general formula (IV) is hydrolyzed steeply when the reaction temperature is too high. In terms of the reaction time, though it varies depending on the amount of the reaction solution, it is preferable to add dropwisely the silane compound represented by the above described general formula (IV) to said solution for from 2 to 6 hours, and then the solution is kept stirred for from 5 to 20 hours after said addition completed.

At completion of the hydrolysis reaction, because the reaction solution is in acidic or basic state, the solution is neutralized by adding an aqueous alkali solution such as an aqueous sodium hydroxide solution, aqueous potassium hydroxide solution and the like when the reaction solution is acidic, or by adding an acid aqueous solution such as dilute hydrochloric acid, dilute sulfuric acid and the like when the reaction solution is basic; and then the solution is stirred to be neutralized. By this neutralization, polycondensation is further proceeded to give a silylated peptide-silane compound copolymer composition. Stirring time conducted after the neutralization is preferably for from 1 to 10 hours.

A silylated peptide-silane compound copolymer

composition having better compatibility in a hair treatment agent containing a large amount of oily material can be produced by suitably selecting the kind and reaction amount of the silane compound represented by the general formula (II) in the polycondensation of the silylated peptide represented by the general formula (I) with the silane compound represented by the general formula (II).

That is, the silylated peptide-silane compound copolymer composition has surface activity, since it is produced by polycondensation of a hydrophilic peptide portion in the silylated peptide and the silane compound having hydrophobic group. As the result, when the hair treatment agent contains a higher fatty acid and a higher alcohol having long carbon chain, stability of the higher fatty acid and the higher alcohol having long carbon chain can be improved, if a silane compound represented by the general formula (IV) and connecting an organic group as R^6 having a longer carbon chain, such as hexyltrimethoxysilane, decyltrimethoxysilane and the like is used to polycondensing with the silylated peptide.

Thus, by selecting kind of the silane compound to be reacted, silane compound copolymer composition having various properties can be obtained.

The silylated peptide-silane compound copolymer composition used for a hair treatment agent, a shampoo or a hair coloring agent of the present invention is produced by polycondensing one or more kind selected from silylated

peptide represented by the above described general formula (I) with one or more kind selected from silane compound represented by the above described general formula (II) preferably in a range of reaction molar ratio of from 1:1 to 1:100, more preferably from 1:1 to 1:85.

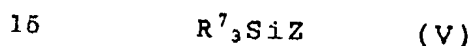
When the reaction molar ratio of the silylated peptide represented by the general formula (I) to the silane compound represented by the general formula (II) is less than the above described range, a hair treatment agent containing said copolymer composition is impossible to provide properties such as imparting gloss and springness to hairs; a shampoo containing said copolymer composition is impossible to provide properties such as imparting gloss to hairs, making hairs smooth and improving combability of hairs; and a hair coloring agent containing said copolymer composition is impossible to provide properties such as giving little damage to hairs on dyeing, imparting good gloss and springness to hairs, and exhibiting good shampoo fastness.

When the reaction molar ratio of the silylated peptide represented by the general formula (I) to the silane compound represented by the general formula (II) is more than the above described range, the hair treatment agent, the shampoo and the hair coloring agent not only is unable to achieve the increase of properties imparting gloss, springness and smoothness to meet the value of the reaction molar ratio but causes silicon-oil touch and further may deteriorate handling ability because of high viscosity of the silylated

peptide-silane compound copolymer composition .

Because a hydroxyl group remains at terminal silylate group of the silylated peptide-silane compound copolymer composition obtained by above described procedure, the silylated peptide-silane compound copolymer compositions may coagulate themselves and, as a result, be highly polymerized. Therefore, a silane compound represented by the above described general formula (III) which produces with hydrolysis one hydroxyl group directly connecting to silicon atom is reacted with said copolymer composition.

The silane compound represented by the above described general formula (III) is obtained by hydrolyzing a silane compound represented by the following general formula (V) in aqueous solution



, wherein three R^7 's represent organic groups in which a carbon atom is directly connected to the silicon atom and three R^7 's may be the same or different, Z is at least one group selected from the group consisting of a hydroxyl group, an alkoxy group and a halogen group.

Examples of the silane compound represented by the general formula (V) which produces with hydrolysis one hydroxyl group directly connecting to the silicon atom, include dimethylvinylchlorosilane, n-butyldimethylchlorosilane, tert-butyldimethylchlorosilane, tert-butyldiphenylchlorosilane, octadecyldimethylchlorosilane,

methyldiphenylchlorosilane, tri-n-butylchlorosilane,
triethylchlorosilane, trimethylchlorosilane,
tri-n-propylchlorosilane, triphenylchlorosilane,
trimethyliodidesilane, trimethylethoxysilane,
5 dimethylvinylethoxysilane, dimethylvinylmethoxysilane,
trimethylethoxysilane, trimethylmethoxysilane,
triphenylethoxysilane and the like.

In addition, silyl compounds having two silicon atoms
such as hexamethyldisilazane and hexamethyldisiloxane can
10 also be used because these compounds generate one hydroxyl
group directly connecting to a silicon atom with hydrolysis.

Since a silane compound represented by the general
formula (V) which produces by hydrolysis one hydroxyl group
directly connecting to silicon atom has only one reactive
15 group directly connected to a silicon atom; a silane compound
represented by the general formula (III) obtained by
hydrolyzing above said silane compound reacts with a
hydroxyl group existing in the silylated peptide-silane
compound copolymer composition; and reduces the hydroxyl
20 group in said silylated peptide-silane compound copolymer
composition; and thus prevents said silylated
peptide-silane compound copolymer composition from further
polycondensing by themselves. Namely, by reacting the
silane compound represented by the general formula (III),
25 a silylated peptide-silane compound copolymer composition
having excellent storage stability can be obtained.

A reaction of a silylated peptide-silane compound
copolymer composition with a silane compound represented

by the general formula (III) is conducted, for example,
by adding dropwisely the silane compound represented by
the general formula (V) into an aqueous solution of the
silylated peptide-silane compound copolymer composition;
5 and thus a hydroxyl group of the silylated peptide-silane
compound copolymer composition is condensed with a hydroxyl
group of the silane compound represented by the general
formula (III).

In case using a silane compound, in which Z represents
10 a halogen atom, as the silane compound represented by the
above described general formula (V), because said silane
compound shows excellent hydrolysis property, the
above-described reaction can be conducted by directly
adding dropwisely the silane compound represented by the
15 above described general formula (V) into an aqueous solution
of the silylated peptide-silane compound copolymer
composition. However, in case using a silane compound, in
which Z represents an alkoxy group or a silane compound
containing two silicon atoms such as hexamethyldisiloxane,
20 as the silane compound represented by the above described
general formula (V), it is necessary that said silane
compound is previously hydrolyzed in an aqueous solution
having pH 2 to 3 to give a silane compound represented by
the general formula (III) and then the silane compound
25 obtained by hydrolysis is added dropwisely into an aqueous
solution of the silylated peptide-silane compound
copolymer composition.

The reaction temperature of an silylated

peptide-silane compound copolymer composition with a silane compound represented by the above described general formula (V) is preferably from 30 to 60 °C. The reaction time, though it varies depending on the amount of reaction solution, is preferably for from 30 minutes to 2 hours to add dropwisely the silane compound represented by the general formula (V), and then for from 1 to 6 hours to stir the reaction solution after said addition completed.

After completion of the stirring, the reaction solution is neutralized with an aqueous alkali solution such as an aqueous sodium hydroxide solution, an aqueous potassium hydroxide solution and the like; and the reaction is completed by further stirring for 1 to 10 hours to obtain a silylated peptide-silane compound copolymer composition. After controlling pH and concentration of the reaction solution, the silylated peptide-silane compound copolymer composition having viscosity in a range from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C is used for a hair treatment agent, a shampoo or a hair coloring agent of the present invention.

The reason why an amount of the silylated peptide-silane compound copolymer composition contained in the hair treatment agent, the shampoo or the hair coloring agent of the present invention is set in the range in terms of the viscosity ranging from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C is; that an effect of silylated peptide-silane compound copolymer composition may not be

realized due to insufficient polymerization when the viscosity in 70% of solid content concentration of said copolymer composition at 20 °C is less than the above described range; and that a handling ability of the silylated peptide-silane compound copolymer composition is
5 deteriorated due to the lack of fluidity when the viscosity is more than the above described range.

The hair treatment agent of the present invention
10 includes a hair rinse, hair conditioner, hair treatment, hair treatment cream, PPT (polypeptide) treatment, hair cream, hair spray, hair wax and the like, a shampoo, a hair coloring agent, and the like, and comprises the above-mentioned a silylated peptide-silane compound
15 copolymer composition.

In general, an amount contained in the hair treatment agent of the present invention (that is, an amount blended to the hair treatment agent) of the silylated peptide-silane compound copolymer composition is preferably from 0.01 to
20 5% by weight in the hair treatment agent. However, the preferable amount varies depending on kind and method of use of the hair treatment agent, since it may cause roughness and stickiness on the hair due to the water-insoluble property of the silylated peptide-silane compound
25 copolymer composition and to accumulation of said copolymer composition in daily use when the amount contained of the silylated peptide-silane compound copolymer composition is large.

Therefore, the preferable amount is 0.01 to 1% by weight in the hair treatment agent for a hair rinse to be used daily; and the preferable amount is 0.05 to 5% by weight for a hair treatment applied to damaged hairs due to a chemical treatment such as a permanent wave treatment or dyeing hair. When a dyed hair treatment to be used after dyeing hairs contains the silylated peptide-silane compound copolymer composition in an amount of 0.05 to 5% by weight, elimination of dye from the dyed hair is suppressed. Regarding any kinds of hair treatment agent, the effects of imparting excellent gloss and springness to hairs, improving combability of hairs, preventing formation of split hairs, improving manageability of hairs, and preventing discoloration of dyed hairs due to shampoo, when used for dyed hairs, may not be attained, when content of the silylated peptide-silane compound copolymer composition is 0.01% by weight or less.

Of a shampoo of the present invention, an amount contained in the shampoo (that is, an amount blended to the shampoo) of the silylated peptide-silane compound copolymer composition having viscosity in a range from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C is preferably from 0.01 to 5% by weight, more preferably from 0.05 to 2.5% by weight. It may be unable to fully obtain the effects imparting gloss to hairs, making hairs smooth, improving combability and manageability of hairs, when the amount contained of the

silylated peptide-silane compound copolymer composition is less than the above described range. It may cause roughness and stickiness on the shampooed hair due to the water-insoluble property of the silylated peptide-silane compound copolymer composition and to accumulation of said copolymer composition in daily use when the amount contained of the silylated peptide-silane compound copolymer composition is more than the above described range. In blending to the shampoo, the silylated peptide-silane compound copolymer composition may be used alone or in a mixture of more than two kinds.

The shampoo of the present invention may be produced with conventional manner except for containing the silylated peptide-silane compound copolymer composition having viscosity in a range from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C.

For example, various surfactants used conventionally may be used for a surfactant used as a major ingredient of a shampoo. For example, anionic surfactants, nonionic surfactants, ampholytic surfactants, cationic surfactants, and the like may be used alone or in a mixture of more than two kinds. An amount of the surfactant used is usually from 0.1 to 25% by weight in the shampoo, more preferably from 2 to 25% by weight, and further more preferably from 5 to 15% by weight.

Examples of the anionic surfactants include alkyl sulfates such as ammonium lauryl sulfate, lauryl

ethanolamine sulfate, sodium lauryl sulfate, lauryl triethanolamine sulfate and the like; polyoxyethylene alkyl ether sulfates such as polyoxyethylene(2EO)laurylether triethanolamine sulfate
5 (,wherein EO represents ethylene oxide and the value attached on the head of letter of EO represents the number of ethylene oxide molecules added), sodium polyoxyethylene(3EO)alkyl(,wherein said alkyl group is selected from the group having carbon atoms from 11 to 15
10 or from a mixture of more than two kinds thereof, and is abbreviated in a manner such as "C11-15" hereinafter)ether sulfate, and the like; alkylbenzene sulfates such as sodium laurylbenzene sulfate, laurylbenzene triethanolamine sulfate and the like; polyoxyethylene alkyl ether acetates
15 such as sodium polyoxyethylene(3EO)tridecylether acetate and the like; sodium alkane sulfate, sodium hydrogenated(palm oil fatty acid)monoglyceride sulfate, disodium undecylenoylamideethyl sulfosuccinate, sodium octylphenoxydiethoxyethyl sulfate, disodium oleamide
20 sulfosuccinate, sodium dioctyl sulfosuccinate, disodium lauryl sulfosuccinate, polyoxyethylenealkyl(C12-16)ether phosphate(2-12 EO), sodium polyoxyethyleneoleylether phosphate, sodium polyoxyethylenecetylether phosphate, disodium polyoxyethylenelaurylether sulfosuccinate,
25 sodium polyoxyethylenelaurylether phosphate, sodium lauryl sulfosuccinate, sodium tetradecene sulfonate and the like.

Examples of nonionic surfactants include

polyoxyethylenealkyl(C12-14) ether(7EO),
 polyoxyethyleneoctylphenyl ether, polyoxyethyleneoleyl
 ether, glyceryl polyoxyethylene oleate,
 polyoxyethylenestearyl ether,
 5 polyoxyethylenecetyl ether, polyoxyethylenecetylstearyl
 diether, polyoxyethylenesorbitol inolin acid(40EO),
 polyoxyethylenenonylphenyl ether,
 polyoxyethylenepolyoxypropylenecetyl ether,
 polyoxyethylenepolyoxypropylenedecyltetradecyl ether,
 10 decylpolyglycoxyde, laurylpolyglycoxyde,
 cetylpolyglycoxyde, polyoxyethylenelanolin alcohol,
 polyoxypropylenestearyl ether and the like.

Examples of ampholytic surfactants include
 2-alkyl-N-carboxylmethyl-N-hydroxyethyl imidazolinium
 15 betaine, undecylhydroxyethyl imidazolinium betaine,
 undecyl-N-hydroxyethyl-N-carboxylmethyl imidazolinium
 betaine, stearyldihydroxyethyl betaine, stearyldiethyl
 betaine, cocoalkyldimethyl glycine,
 cocoamidopropyldimethyl glycine, sodium
 20 cocoalkyl-N-carboxyethyl-N-hydroxyethyl imidazolinium
 betaine, disodium
 cocoalkyl-N-carboxyethoxyethyl-N-carboxyethyl
 imidazolinium hydroxide, DL-pyrrolidonecarboxylic acid
 salt of L-cocoylarginine ethyl ester
 25 and the like.

Examples of cationic surfactants include
 stearyltrimethyl ammonium chloride, cetyltrimethyl
 ammonium chloride, cetostearyltrimethyl ammonium chloride,

stearylbis(diethyleneglycol)hydroxyethyl ammonium chloride, behenyltrimethyl ammonium chloride, distearyldimethyl ammonium bromide, stearyldimethylbenzyl ammonium chloride,

5 behenyltrimethyl ammonium bromide, cetyltrimethyl ammonium iodide, oleylbenzyldimethyl ammonium chloride, oleylbis[polyoxyethylene(15EO)]methyl ammonium chloride, tri[polyoxyethylene(5EO)]stearyl ammonium chloride, alkyltrimethyl ammonium chloride,
10 polyoxypropylenemethyldiethyl ammonium chloride, mink oil amidepropyldimethylhydroxyethyl ammonium chloride, alkylpyridinium salt, γ -gluconamidepropyldimethylhydroxyethyl ammonium chloride, and the like.

15

An amount contained in the hair coloring agent (that is, an amount blended to the hair coloring agent) of the silylated peptide-silane compound copolymer composition is preferably from 0.05 to 10% by weight, particularly
20 preferably from 0.5 to 5% by weight. It may be unable to fully obtain the effects suppressing damage of hairs on dyeing and imparting gloss, springness and good combability to hairs, and shampoo fastness of dyed hairs, when the amount contained of the silylated peptide-silane compound
25 copolymer composition is less than the above described range. Improvement of the effects meeting the increase of the amount scarcely realizes, and it may cause stickiness on the hair and oily feeling due to a large amount of the silylated

peptide-silane compound copolymer composition remained on the hair, when the amount contained of the silylated peptide-silane compound copolymer composition is more than the above described range.

5

For a two-package type hair coloring agent, one embodiment of the present invention, the silylated peptide-silane compound copolymer composition may be comprised either in a first composition or in a second
10 composition. In general, it is more suitable to be comprised in the first composition. In general, the first composition of a two-package type hair coloring agent comprises an oxidation dye (dye intermediate) and an alkali agent, and further, based on demand, a nitro dye.

15 Examples of the oxidation dye include a phenylenediamine compound such as p-phenylenediamine, N-phenyl-p-phenylenediamine, 4,4'-diaminodiphenylamine or o-phenylenediamine, a tolueneamine compound such as a toluene-2,5-diamine or toluene-3,4-diamine, an
20 aminophenol compound such as p-aminophenol, p-methylaminophenol or o-methylaminophenol, an aminonitrophenol such as o-amino-m-nitrophenol, a diaminopyridine such as 2,6-diaminopyridine, and the like. Generally used couplers include, for example,
25 m-phenylenediamine, toluene-2,4-diamine, m-aminophenol, naphthol, resorcinol, catechol, hydroquinone and the like.

The alkali agent is not particularly limited, and examples thereof include sodium hydroxide, potassium

hydroxide, aqueous ammonia, monoethanol amine, triethanol amine, and the like. Examples of nitro dye include 2-amino-4-nitrophenol, nitro-p-phenylenediamine, p-nitro-o-phenylenediamine, 2-amino-5-nitrophenol, and
5 the like.

The second composition contains an oxidizing agent as the essential component, and the oxidizing agent is not particularly limited. Examples of the oxidizing agent used in the second composition include hydrogen peroxide, sodium
10 percarbonate, sodium perborate, sodium peroxide, and the like.

Hair coloring agent of the present invention comprises a silylated peptide-silane compound copolymer composition having viscosity in a range from 500 to 20000 mPa·s in
15 70% of solid content concentration of said copolymer composition at 20 °C, as the essential component, and also comprises another component as long as it is usable in the hair coloring agent. Preparation of the hair coloring agent of the present invention can be conducted either by
20 adding a silylated peptide-silane compound copolymer composition having viscosity in a range from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C to an already prepared hair coloring agent, or by adding a silylated peptide-silane compound
25 copolymer composition having viscosity in a range from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C together with other component simultaneously.

As mentioned above, a hair treatment agent of the present invention essentially can be prepared by adding a silylated peptide-silane compound copolymer composition
5 having viscosity in a range from 500 to 20000 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C to a conventional hair treatment agent. Furthermore, other suitable components may be added to the hair treatment agent of the present invention in the range
10 not to deteriorate the effects of the present invention.

Example of the other suitable component include anionic surfactants, nonionic surfactants, cationic surfactants, ampholytic surfactants, cationic polymers, ampholytic polymers, anionic polymers, thickener, extracts
15 from animals and vegetables, polysaccharide or derivatives thereof, hydrolyzed peptides of proteins derived from animals, vegetables and microorganism or derivatives thereof, wetting agent, lower alcohols, higher alcohols, amino acids, fats and oils, silicones, preservatives,
20 perfumes, and the like.

A shampoo of the present invention essentially comprises a silylated peptide-silane compound copolymer composition having viscosity in a range from 500 to 20000
25 mPa·s in 70% of solid content concentration of said copolymer composition at 20 °C and a surfactant, and said shampoo is produced by dissolving said those essential components into water or a solution comprising water and a suitable solvent.

Furthermore, other suitable components may be added in the range not to deteriorate the effects of the present invention.

Example of said other suitable components are
5 illustrated with as follows:

A synthetic polymer such as cationic polymer, ampholytic polymer and anionic polymer including cationized cellulose, cationized guar gum, polydiallyldimethyl ammonium chrolide, polyvinylpyrrolidone, polyethyleneimine and the like; A
10 thickeners such as isostearic acid diethanolamide, undecylic acid monoethanolamide, oleic acid diethanolamide, monoethanolamine tallow acid amide, hydrogenated tallow acid diethanolamide, stearic acid diethanolamide, stearic acid diethylaminoethylamide, stearic acid
15 monoethanolamide, myristic acid diethanolamide, coconut fatty acid ethanolamide, coconut fatty acid diethanolamide, lauric acid isopropanolamide, lauric acid ethanolamide, lauric acid diethanolamide, lanoline fatty acid diethanolamide, and the like; a fat and an oil such as waxes,
20 paraffin, fatty acid esters, glyceride, oils derived from animals and vegetables, and the like; a moisturizing agent such as a substance extracted from animals and vegetables, polysaccharide or its derivatives, propyleneglycol, 1,3-butyleneglycol, ethyleneglycol, glycerin, and the
25 like; a lower alcohol such as ethanol, propanol, isopropanol, and the like; a higher alcohol such as cetanol, stearyl alcohol, oleyl alcohol, behenyl alcohol, and the like; an amino acid such as L-aspartic acid, sodium L-aspartate,

DL-alanine, L-arginine, glycine, L-glutamic acid,
L-cystine, L-threonine, and the like; a licorice derivative
such as glycyrrhizinic acid, disodium carbenoxolone, and
the like; an antiphlogistic such as allantoin, guaiazulene,
5 aloe, a-bisabolol, and the like; an anti-dandruff agent
such as salicylic acid, zinc pyrithione, pyroctone olamine,
sulfur, selenium sulfide, triclosan, resorcin, vitamin A,
vitamin B₆, and the like; a hydrolyzed protein derived from
animal, vegetable and microorganism such as soy bean, wheat,
10 corn, potato, yeast, fungi and the like, and a derivative
such as N-quaternary ammonium derivatives,
N-acylated derivatives, N-silylated derivatives,
esterified derivatives, and the like, wherein said animal-,
vegetable- and microorganism-derived protein include
15 collagen, keratin, silk, serine, casein, and the like; a
silicon derivative such as chained or cyclic
methylpolysiloxane, methylphenylpolysiloxane,
dimethylpolysiloxane-polyethyleneglycol copolymer, amino
modified silicon, quaternary ammonium modified silicon,
20 and the like.

A hair coloring agent of the present invention may
further comprise other suitable components in the range
not to deteriorate the effects of the present invention,
25 as conventional hair coloring agents, in addition to the
above-mentioned essential components.

Example of the other suitable component include
surfactants such as nonionic surfactants, anionic

surfactants, cationic surfactants, ampholytic surfactants,
a dissolving assistant such as glycerol or propylene glycol,
a humectant, a viscosity modifier such as
carboxymethylcellulose, hydroxyethylcellulose, carboxy
6 vinylpolymer or polyacryl amide, copolymer of acrylates,
methacrylates or the like, polymers such as
N,N-dimethylaminoethylmethacrylate copolymer, copolymer
of vinylpyrrolidone and vinyl acetate or the like, pH
adjusting agents, perfumes, presevatives, antioxidants,
10 chelating agents, and the like.

The hair treatment agent of the present invention
imparts good gloss and springness to hairs, improves
combability of hairs, prevents formation of split hairs,
15 improves manageability of hairs, and particularly, is
excellent in a property which prevents discoloration of
hairs due to shampoo, when used for dyed hairs, due to the
silylated peptide-silane compound copolymer composition
having viscosity in a range from 500 to 20000 mPa·s in
20 70% of solid content concentration of said copolymer
composition at 20 °C contained in the composition.

The shampoo of the present invention imparts gloss
to hairs, makes hairs smooth, improves combability and
25 manageability of hairs.

The hair coloring agent of the present invention dyes
hairs uniformly, gives little damage to hairs on dyeing,

and imparts good gloss, springness, combability and the like to hairs, and makes shampoo fastness of dyed hairs excellent.

5 [Examples]

The present invention is more specifically described and explained by means of the following Examples, but is not limited the scope thereof. In the following Example and Comparative Example, the amount of each component
10 blended is represented by the part by weight; and if the amount blended is not the amount of solid content of said component, the concentration of said solid content is represented by being enclosed in parentheses followed to said component name. The percent (%) showing concentration
15 is % by weight. In some Examples and Comparative Examples, a term "blend" is used in stead of "contain". Prior to the Example, examples producing a silylated peptide-silane compound copolymer composition used in the Example are described as Production Example.

20

Production Example 1

Production of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxylpropyl
hydrolyzed

25 sericin-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition [1:25:25 (molar ratio)]

Into the reaction vessel made of glass shaped in cylinder with round bottom of its inner diameter of 12 cm

and its volume of 2 liters, 200 g of 10% aqueous solution of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed sericin (, wherein the number average molecular

5 weight of said hydrolyzed sericin is about 500) and 11.5 g of 18% aqueous solution of hydrochloric acid were added

to adjust pH of a mixture to 1.5 and said mixture was heated up to 60 °C. While the mixture being stirred in 400 rpm,

10 the mixture of 99.1 g of dimethyldiethoxysilane (a trade name of KBE-22, manufactured by Shin-Etsu Silicone Co.,

Ltd.) and 184.7 g of octyltriethoxysilane (a trade name of A-137, Nippon Unicar Co., Ltd.) was dropped into said

mixture for 5 hours, and then, after said dropping being completed, the solution obtained was further stirred for

15 15 hours at 60 °C. Then, while the solution being stirred, 40.8 g of 5% aqueous solution of sodium hydroxide was

gradually dropped into said solution to adjust the solution pH to 6, and the solution was further stirred for one hour

at 60 °C. While the reaction solution obtained being stirred

20 in 400 rpm at 60 °C, 11.6 g of trimethylchlorosilane (a trade name of KA-31, manufactured by Shin-Etsu Silicone Co., Ltd.)

was added, and then said reaction solution was stirred for one hour at 60 °C. Then, after 78.8 g of 5% aqueous solution

of sodium hydroxide being dropped into the reaction solution

25 and pH of said solution being adjusted to 6, the solution was stirred for one hour at 60 °C, and then the solution

was heated up to 80 °C and further stirred for one hour.

Then, the reaction solution was concentrated to the solid

content of 70% by vacuum concentration with the rotary-evaporator.

Thus, 260 g of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
5 hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition was obtained.

The viscosity of 70% aqueous solution at 20 °C of
obtained

10 N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition was measured by B-type viscometer
with the rotor 3 and number of revolution 30, thus the
15 viscosity resulted in 2120 mPa·s.

Production Example 2

Production of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
20 hydrolyzed

collagen-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition [1:25:25 (molar ratio)]

Into the reaction vessel made of glass shaped in
cylinder with round bottom of its inner diameter of 12 cm
25 and its volume of 2 liters, 150 g of 10% aqueous solution
of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed collagen (, wherein the number average molecular

weight of hydrolyzed collagen is about 500) and 7.6 g of 18% aqueous solution of hydrochloric acid were added to adjust pH of the mixture to 1.5 and said mixture was heated up to 60 °C. While the mixture being stirred in 400 rpm, the mixture of 79.7 g of dimethyldiethoxysilane (a trade name of KBE-22, manufactured by Shin-Etsu Silicone Co., Ltd.) and 148.6 g of octyltriethoxysilane (a trade name of A-137, Nippon Unicar Co., Ltd.) was dropped into said mixture for 5 hours, and then, after said dropping being completed, the solution obtained was further stirred for 15 hours at 60 °C. Then, while the solution being stirred, 22.9 g of 5% aqueous solution of sodium hydroxide was gradually dropped into said solution to adjust the solution pH to 6, and the solution was further stirred for one hour at 60 °C. While the reaction solution obtained being stirred in 400 rpm at 60 °C, 9.3 g of trimethylchlorosilane (a trade name of KA-31, manufactured by Shin-Etsu Silicon Co., Ltd.) was added, and then said reaction solution was stirred for one hour at 60 °C. Then, after 68.5 g of 5% aqueous solution of sodium hydroxide being dropped into the reaction solution and pH of said solution being adjusted to 6, the solution was stirred for one hour at 60 °C, and then the solution was heated up to 80 °C and further stirred for one hour. Then, the reaction solution was concentrated to the solid content of 70% by vacuum concentration with the rotary-evaporator.

Thus, 211.9 g of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl

hydrolyzed

collagen-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition was obtained.

5 The viscosity of 70% aqueous solution at 20 °C of
obtained

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed

collagen-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition was measured according to the same
10 condition applied in Production Example 1, thus the
viscosity resulted in 1116 mPa·s.

Production Example 3

Production of

15 N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed

silk-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition [1:40:40 (molar ratio)]

20 Into the reaction vessel made of glass shaped in
cylinder with round bottom of its inner diameter of 12 cm
and its volume of 2 liters, 127.3 g of 10% aqueous solution
of

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed silk (, wherein the number average molecular
25 weight of hydrolyzed silk is about 600) and 4.8 g of 18%
aqueous solution of hydrochloric acid were added to adjust
pH of the mixture to 1.5 and said mixture was heated up
to 60 °C. While the mixture being stirred in 400 rpm, the

mixture of 88.0 g of dimethyldiethoxysilane (a trade name of KBE-22, manufactured by Shin-Etsu Silicon Co., Ltd.) and 164.0 g of octyltriethoxysilane (a trade name of A-137, Nippon Unicar Co., Ltd.) was dropped into said mixture for 5 and half hours, and then, after said dropping being completed, the solution obtained was further stirred for 15 hours at 60 °C. Then, while the solution being stirred, 17.1 g of 5% aqueous solution of sodium hydroxide was gradually dropped into said solution to adjust the solution pH to 6, and the solution was further stirred for one hour at 60 °C. While the reaction solution obtained being stirred in 400 rpm at 60 °C, 6.4 g of trimethylchlorosilane (a trade name of KA-31, manufactured by Shin-Etsu Silicon Co., Ltd.) was added, and then said reaction solution was stirred for one hour at 60 °C. Then, after 45.7 g of 5% aqueous solution of sodium hydroxide being dropped into the reaction solution and pH of said solution being adjusted to 6, the solution was stirred for one hour at 60 °C, and then the solution was heated up to 80 °C and further stirred for one hour. Then, the reaction solution was concentrated to the solid content of 70% by vacuum concentration with the rotary-evaporator.

Thus, 209 g of N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxylpropyl hydrolyzed silk-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition was obtained.

The viscosity of 70% aqueous solution at 20 °C of

obtained

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed

silk-dimethyldiethoxysilane-octyltriethoxysilane

5 copolymer composition was measured by the same condition
applied in Production Example 1, thus the viscosity resulted
in 640 mPa · s.

Example 1 and Comparative Examples 1 and 2

10 Three kinds of hair treatment agents shown in Table
1 were prepared. After applying the hair treatment agents
on dyed hairs respectively, gloss, springness, combability
and manageability of the treated hairs were evaluated. In
addition, degree of discoloration of the treated hairs by
15 shampooing was also visually compared with naked eyes.

In Example 1,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane

20 copolymer composition produced in Production Example 1,
using a silylated peptide-silane compound copolymer
composition, was used. In Comparative Example 1, in place
of a silylated peptide-silane compound copolymer
composition, polyoxyethylene/methylpolysiloxane
25 copolymer, a polyether modified silicone, was used; in
Comparative Example 2, neither a silylated peptide-silane
compound copolymer composition nor a silicone were used.

The numbers representing amounts in all Tables in Examples and Comparative examples in this specification are weight parts unless otherwise stated.

5 Table 1

	Example	Comparative example	
	1	1	2
Silylated peptide-silane compound copolymer composition (70%) of Production Example 1	3.0	0	0
Polyoxyethylene/methylpoly siloxane copolymer*1	0	3.0	0
Sodium acrylate/acryloyldimethyl Taurine copolymer +isohexadecane+ Polysorbate80 mixture *2	2.0	2.0	2.0
1,3-butleneglycol	2.0	2.0	2.0
glycerol	1.0	1.0	1.0
Liquid paraffin#70	1.0	1.0	1.0
Squalane	2.0	2.0	2.0
Hydrolyzed silk (7%) *3	0.5	0.5	0.5
Paraoxybenzoate+phenoxy ethanol mixture *4	0.5	0.5	0.5
Sterilized ion-exchange water	Volume adjust-able totally 100 part	Volume adjust-able totally 100 part	Volume adjust-able totally 100 part

*1: SH3775M (trade name), manufactured by Dow Corning Toray Silicone Co.,Ltd.

*2: Simul gel EG (trade name), manufactured by Seppic

Co., Ltd.

*3: Promois silk-1000(trade name), manufactured by SEIWA KASEI Co., Ltd.

*4: Seisept H(trade name, manufactured by SEIWA KASEI Co.,
5 Ltd.

Dyed hair samples to be treated for evaluation were prepared by washing, bleaching and dyeing hairs.

More concrete steps are as follows:

- 10 Preparing three bundles of hair, each bundle being the length of 13 cm and the weight of 1.7 g;
Washing a bundle with 2% aqueous solution of sodium polyoxyethylene(3) laurylether sulfate, and then rinsing it with streaming tap water; and then
- 15 Drying it in the air to provide a hair bundle;
Dipping said hair bundle for 30 minutes in a bleaching agent, a mixture of 6% aqueous hydroperoxide and 2% aqueous ammonia in the weight ratio of 1:1,
Rinsing it with streaming tap water,
- 20 Dipping said hair bundle for 5 minutes in a buffer solution of 1 M citric acid and 0.2 M disodium hydrogenphosphate (pH3),
Rinsing it with streaming tap water,
Drying it at room temperature in the air,
- 25 Repeating above procedure 5 times,
Dyeing hair with a commercially available oxidation type hair coloring agent for black,
Getting the dye hair bundle wet with hot water at 40 °C,

Coating 3 g of the hair treatment agent obtained in Example 1 or Comparative examples 1 or 2 on the hair bundle and spreading and permeating them thoroughly,
Leaving it on a wrap film in a thermostat of 40 °C for 10
5 minutes,

Rinsing it with hot water, and

Drying it at room temperature in the air.

By using the hair bundles thus prepared, the properties
10 of gloss, springness, combability and manageability of hair were sensory evaluated by 10 sensory-panelists in rating the properties as follows: best (point 2), second best (point 1) and worse (point 0); and then the average of rated points was defined as the evaluation value.

15

Then, the bundles of hair evaluated above were dipped in 200 ml of 1% (effective content) aqueous sodium polyoxyethylene(3) laurylether sulfate for 10 minutes, rinsed with streaming tap water, and then dried at room
20 temperature in the air. After this procedure was repeated 10 times, the dyeing power of dyed hair were visually evaluated by 10 sensory-panelists in rating the properties as follows: deepest, (point 2), second deepest (point 1) and faint (point 0); and then the average of rated points
25 was defined as the evaluation value for discoloration of dyed hair.

The results of evaluations are shown in Table 2 in terms of averaged value of ten panelists.

Table 2

	Example	Comparative example	
	1	1	2
Treated hair:			
Gloss	1.8	1.0	0.2
Springness	1.7	0.8	0.5
Combability	1.9	0.7	0.4
Manageability	1.5	1.0	0.5
Depth of color of dyed hair	1.8	0.7	0.5

As shown in Table 2, the evaluated values of the hair
5 treated by the hair treatment agent of Example 1 are higher
in every evaluation item such as gloss, springness,
combability and manageability of hair in comparison with
the hairs treated by the hair treatment agent of Comparative
Examples 1 and 2. Therefore, the hair treatment agent of
10 Example 1 blended with
N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxylpropyl
hydrolyzed
sericin-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition clearly exerts excellent effects
15 imparting gloss and springness to hairs, improving
combability and manageability of hairs than the hair
treatment agent of Comparative example 1 blended with
polyoxyethylene/methylpolysiloxane copolymer and of
Comparative example 2 not blended with a silylated
20 peptide-silane compound, etc.

In the comparison of color depth of the dyed hair, almost all panelists answered that the color of hair treated with the hair treatment agent of Example 1 was deepest.

5 This result shows that the hair treatment agent of Example 1 blended with

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane

10 copolymer composition clearly exerts excellent effects suppressing discoloration by shampooing dyed hairs.

Example 2 and Comparative Example 3

Two kinds of hair creams shown in Table 3 were prepared.

15 After applying them on dyed hairs respectively, gloss, springness, combability and manageability of the treated hairs were evaluated.

In Example 2,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl

20 hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition produced in Production Example 1, using a silylated peptide-silane compound copolymer composition, was used. In Comparative Example 3, in place

25 of a silylated peptide-silane compound copolymer composition,

poly(oxyethylene/oxipropylene)/methylpolysiloxane copolymer, a polyether modified silicone, was used.

Table 3

	Example	Comparative example
	2	3
Silylated peptide-silane compound copolymer composition (70%) of Production Example 1	0.1	0
Poly(oxyethylene/oxipropylene)/methylpolysiloxane copolymer*5	0	0.1
Paraffin wax	10.0	10.0
Cetanol	2.5	2.5
Stearic acid	4.5	4.5
Glyceryl monoisostearate *6	0.5	0.5
2-ethylhexyl palmitate*7	15.0	15.0
Liquid paraffin#350	15.0	15.0
Triethanol amine	1.8	1.8
Propylene glycol	1.0	1.0
(Dihydroxymethylsilyl propoxy)hydroxyl propyl hydrolyzed wheat protein (20%) *8	0.1	0.1
Presevative	Appropriate amount	Appropriate amount
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part

*5: SH3749 (trade name), manufactured by Dow Corning Toray
 5 Silicone Co., Ltd.

*6: Ayacol GMIS (trade name), manufactured by SEIWA KASEI

Co., Ltd.

*7: Matlube 2EHP (trade name), manufactured by SEIWA KASEI Co., Ltd.

*8: Promois WG-SIG (trade name), manufactured by SEIWA KASEI Co., Ltd.

Ten female panelists treated their hairs, each of which had been treated at least once by any of chemical treatments selected from bleaching, dyeing and permanent waving, with the hair cream of Comparative example 3 for the first 7 days and with the hair cream of Example 2 for the next 7 days, and comparison was carried out.

The treatment of hair with the cream was carried out by putting an appropriate amount (which varies depending on the amount of hairs) of hair cream on the palm of a hand, coating the cream with hand on hair, particularly on severely damaged hair tip, as a manner to rub the cream in the hair, and drying it with a hair dryer.

After the 14 days (that is, after completion of 7 days for use of the cream of Example 2), they answered, regarding gloss, springness, combability and manageability of hair, which of creams of Example 2 and Comparative Example 3 is better or both are not different (, or almost same).

The results are shown in Table 4 in the numbers of the panelists who answered as Example 2 being better, who answered as Comparative Example 3 being better and who answered as there being no difference.

Table 4

	Number of persons who answered "Example 2 is better"	Number of persons who answered "Comparative example 3 is better"	Number of persons who answered "Cannot tell difference"
Treated hair:			
Gloss	8	2	0
Springness	8	2	0
Combability	9	1	0
Manageability	7	3	0

As shown in Table 4, the largest number of panelists answered that the hair cream of Example 2 blended with

5 N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed sericin-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition is better than the hair cream of Comparative example 3 blended with

10 poly(oxyethylene/oxipropylene)/methylpolysiloxane copolymer in every evaluation item. Particularly, almost all panelists who answered "Example 2 is better." told that unnatural force becomes not to be necessary on brushing due to the improvement of combability of hair tip. This

15 indicates that

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed sericin-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition gives effect preventing formation

20 of split hair. These results clearly show that

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition has excellent effect improving gloss,
springness, combability and manageability to used hairs and
preventing formation of split hair from damaged hair.

5

Example 3 and Comparative Examples 4 and 5

Three kinds of hair rinses shown in Table 5 were
prepared. After applying them on damaged hairs respectively,
and gloss, springness, smoothness, combability and
10 manageability of the hairs were evaluated.

In Example 3,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed

collagen-dimethyldiethoxysilane-octyltriethoxysilane
15 copolymer composition produced in Production Example 2 was
used. In Comparative Example 4, in place of a silylated
peptide-silane compound copolymer composition,
polyoxyethylene/methylpolysiloxane copolymer, a
polyether modified silicone, was used. In Comparative
20 Example 5, neither a silylated peptide-silane compound
copolymer composition nor a silicone were used.

Table 5

	Example	Comparative example	
	3	4	5
Silylated peptide-silane compound copolymer composition (70%) of Production Example 2	0.5	0	0
Polyoxyethylene/methyl polysiloxane copolymer*9	0	0.5	0
Diethyl aminoethylamide stearate *10	3.0	3.0	3.0
Ethylene glycol distearate	4.0	4.0	4.0
Diglyceryl monoisostearate *11	1.5	1.5	1.5
Lipophilic glyceryl monostearate	0.5	0.5	0.5
Behenyl alcohol	6.0	6.0	6.0
Cetyl trimethyl ammonium chloride(29%)	3.5	3.5	3.5
Distearyl dimethyl ammonium chloride(75%)	1.5	1.5	1.5
Propylene glycol	3.0	3.0	3.0
Piroctone olamine*12	0.3	0.3	0.3
Hydrolyzed collagen (30%) *13	0.2	0.2	0.2
Edetate disodium	Appropriate amount	Appropriate amount	Appropriate amount
Presevative	Appropriate amount	Appropriate amount	Appropriate amount
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part	Volume adjustable totally 100 part

*9 SH3775 M (trade name), manufactured by Dow Corning Toray Silicone Co., Ltd.

5 *10: Ayacol amineamide 50E (trade name), manufactured by SEIWA KASEI Co., Ltd.

*11: Ayacol GMIS (trade name), manufactured by SEIWA KASEI Co., Ltd.

*12: Octopirox (trade name), manufactured by Lion Co., Ltd.

*13: Promois WU-32R (trade name), manufactured by SEIWA
5 KASEI Co., Ltd.

Three bundles of damaged hairs subjected to bleaching according to the same procedure as in Example 1 were prepared and the following hair rinse treatment was carried out.

10 More concrete steps are as follows:

Washing one of the bundle of damaged hair with 4% aqueous solution of sodium polyoxyethylene(3) lauryl ether sulfate, and then rinsing it with streaming tap water; and then Wiping out water lightly in the bundle with a towel,

15 Coating 2 g of the hair rinse obtained in Example 3 or Comparative examples 4 or 5 on the hair bundle and spreading and permeating them thoroughly,

Leaving it on a wrap film for 5 minutes,

Rinsing it with hot water,

20 Drying it at room temperature in the air, and

Repeating above procedure 3 times.

Thus obtained hair samples were subjected to the following evaluation.

The hair smoothness of the treated hair was evaluated
25 by measuring the friction force of the treated hair surface with the friction tester (a trade name of KES-SE, made by Katotech Co., Ltd.) under the conditions of humidity of $40 \pm 1\%$ and temperature $22 \pm 1^\circ\text{C}$.

The smoothness (, or roughness) is represented in said tester with the value of average deviation of friction coefficient sensed by a friction sensor traversing the specified distance on the sample surface, wherein the unit of said value is dimensionless and the value indicates "more smoothness" when the value becomes smaller. In the present test, measurement was carried out 10 times per one sample and the average of obtained values was defined as the measurement value.

Then, by using the same bundles of hair, the properties of gloss, springness, smoothness, combability and manageability of hair were sensory evaluated by 10 panelists according to the same rating as in Example 1.

The results of evaluations are shown in Table 6 in terms of averaged value of ten panelists.

Table 6

	Example	Comparative example	
	3	4	5
Hair smoothness (The average deviation of friction coefficient)	0.0038	0.0045	0.0048
Treated hair:			
Gloss	1.7	0.9	0.4
springness	1.6	0.8	0.6
Smoothness	1.6	1.0	0.4
Combability	1.9	0.7	0.4
Manageability	1.5	0.8	0.7

As shown in Table 6, the average deviation value of friction coefficient, which indicates hair smoothness, of

the hair treated by the hair rinse of Example 3 is the smallest; and said value is 84.4% of that of the hair treated by the hair rinse of Comparative example 4, and 79.2% of that of the hair treated by the hair rinse of Comparative
5 Example 5.

According to this result,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed
collagen-dimethyldiethoxysilane-octyltriethoxysilane
10 copolymer composition, which is blended in Example 3,
clearly exerts effects to be sorbed on and makes smooth
the hair surface.

Moreover, according to the sensory evaluation, the
evaluated values of the hair treated by the hair rinse of
15 Example 3 are higher in every evaluation item such as gloss,
springness, smoothness, combability and manageability of
hair in comparison with the hairs treated by the hair rinse
of Comparative Example 4 and 5. Therefore,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
20 hydrolyzed
collagen-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition clearly exerts excellent effects
imparting gloss and springness to hairs, improving
smoothness, combability and manageability of hairs.

25

Example 4 and Comparative Example 6

Two kinds of hair conditioners shown in Table 7 were
prepared. After applying them on dyed hairs respectively,

and gloss, springness, smoothness, combability, manageability of the hairs and degree of discoloration of dyed hairs were evaluated.

In Example 4,

- 5 N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed
silk-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition produced in Production Example 3 was
used. In Comparative Example 6, in place of a silylated
10 peptide-silane compound copolymer composition,
polyoxyethylene/methylpolysiloxane copolymer, a
polyether modified silicone, was used.

Table 7

	Example	Comparative example
	4	6
Silylated peptide-silane compound copolymer composition (70%) of Production Example 3	1.0	0
Polyoxyethylene/methylpoly siloxane copolymer*14	0	1.0
Decylalcohol-polyethyleneglycol-polyurethane copolymer/propylene glycol/water mixture*15	2.0	2.0
Behenyl alcohol	2.0	2.0
Stearyl alcohol	1.5	1.5
Isopropyl isostearate*16	3.5	3.5
Lipophilic glyceryl monostearate	0.8	0.8
Diethylaminoethylamide stearate*17	1.0	1.0
Polyoxyethylene (200) monostearate	1.0	1.0
Cocodimonium hydroxypropyl hydrolyzed Keratin (30%)*18	0.1	0.1
Citric acid	0.1	0.1
Preseervative	Appropriate amount	Appropriate amount
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part

*14 SH3775 M (trade name), manufactured by Dow Corning Toray Silicone Co., Ltd.

5 *15: Aculyn 44 (trade name), manufactured by Roam & Haas Co., Ltd.

*16: Matlube I · I (trade name), manufactured by SEIWA KASEI

Co., Ltd.

*17: Ayacol amineamide 50E(trade name), manufactured by
SEIWA KASEI Co., Ltd.

*18: Promois WK-HCAQ (trade name), manufactured by SEIWA
5 KASEI Co., Ltd.

Two bundles of damaged hairs subjected to bleaching
5 times according to the same procedure as in Example 1,
followed by dyed were prepared and, then, the following
10 hair conditioning treatment was carried out.

More concrete steps are as follows:

Washing one of the bundle of damaged hair with 4% aqueous
solution of polyoxyethylene(3)lauryleter sodium sulfate,
and then rinsing it with streaming tap water; and then
15 Wiping out water lightly in the bundle with a towel,
Coating 2 g of the hair conditioner obtained in Example
4 or Comparative example 6 on the hair bundle and spreading
and permeating them thoroughly,
Leaving it on a wrap film for 5 minutes,
20 Rinsing it with hot water,
Drying it at room temperature in the air, and
Repeating above procedure 10 times.

Thus obtained hair samples were sensory evaluated by
25 10 panelists regarding gloss, springness, smoothness,
combability and manageability of hair, and degree of
discoloration of hair, and they answered which of hair
conditioner of Example 4 and Comparative Example 6 is better

or both are not different (, or almost same).

The results are shown in Table 8 in the numbers of the panelists who answered as Example 4 being better, who answered as Comparative Example 6 being better and who
6 answered as there being no difference.

Table 8

	Number of persons who answered "Example 4 is better"	Number of persons who answered "Comparative example 6 is better"	Number of persons who answered "Cannot tell difference"
Treated hair:			
Gloss			
Springness	8	0	2
Smoothness	7	0	3
Combability	6	0	4
Manageability	9	0	1
Discoloration	7	0	3
	9	0	1

As shown in Table 8, the evaluated values of the hair
10 treated by the hair conditioner of Example 4 are higher in every evaluation item of gloss, springness, smoothness, combability and manageability of hair in comparison with the hair treated by the hair conditioner of Comparative Example 6. Regarding evaluation of discoloration, almost
15 all panelists answered that hair treated in Example 4 had deeper color than hair treated in Comparative example 6. Therefore,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed

20 silk-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition clearly exerts excellent effects

imparting gloss and springness to hairs, improving smoothness, combability and manageability of hairs and suppressing discoloration of dyed hair by shampooing.

5 Example 5 and Comparative Example 7

A hair wax was prepared using
N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed
silk-dimethyldiethoxysilane-octyltriethoxysilane
10 copolymer composition produced in Production Example 3.

Table 9-1

	Example 5	Comparative Example 7
Silylated peptide-silane compound copolymer composition (70%) of Production Example 3	0.05	-
Candelilla wax	5.0	5.0
Paraffin wax	15.0	15.0
Polyoxyethylene(10) oleylether	3.0	3.0
Cetanol	0.5	0.5
Liquid paraffin	10.0	10.0
Squalane	10.0	10.0
1,3-Butyleneglycol	3.0	3.0
Sodium hydroxide	Amount to adjust pH6.8	Amount to adjust pH6.8
Sterilized ion-exchange water	Volume adjust- able totally 100 part	Volume adjust- able totally 100 part

Ten panelists used the two waxes obtained above on their hair, and, then, sensory evaluated about gloss, springness, and manageability.

The sensory evaluation test was carried by the half-head method, in which the panelists combed their hair in two sides, that is, right and left sides, and then treated one of sides with one of the two waxes and the rest side with the other wax.

Treatment with the hair wax was carried out by coating an appropriate amount (which varies depending on the amount

and length of hairs) of hair wax on the hair, and rubbing and spreading the wax in whole hair.

Gloss, springness, combability and manageability of hair were sensory evaluated by 10 panelists in rating the properties as follows: better (point 1) and worse (point 0); and then total of rated points by 10 panelists was defined as the evaluation value and shown in Table 9-2.

Table 9-2

	Example 5	Comparative example 7
Treated hair:		
Gloss	10	0
Springness	8	2
Manageability	9	1

As shown in Table 9-2, the evaluated values of the hair treated by the hair wax of Example 5 are higher in every evaluation item of gloss, springness, and manageability of hair in comparison with the hair treated by the hair wax of Comparative Example 7. Therefore, N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed silk-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition contained in a hair wax clearly exerts excellent effects imparting gloss and springness to hairs, and improving manageability of hairs.

Example 6 and Comparative Example 8

A hair spray undiluted solution was prepared using N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed collagen-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition produced in Production Example 2, and the prepared undiluted solution was mixed with dimethylether in a weight ratio of 50:50 to prepare hair spray.

Table 10-1

	Example 6	Comparative Example 8
Silylated peptide-silane compound copolymer composition (70%) of Production Example 2	0.7	-
Acrylic resin alkanolamine solution (50%)	1.0	1.0
Cetanol	0.1	0.1
Methyl polysiloxane*19	0.5	0.5
Polyoxyethylene(25)octylether	0.8	0.8
perfume	Appropriate amount	Appropriate amount
Ethanol	Volume adjustable totally 100 part	Volume adjustable totally 100 part

*19 SH200C-10cs (trade name), manufactured by Dow Corning Toray Silicone Co., Ltd.

Ten panelists sprayed uniformly the hair sprays obtained above on their hair, and, then, evaluated about

gloss, springness, and manageability according to the same rating as in Example 5; and then total of rated points by 10 panelists was defined as the evaluation value and shown in Table 10-2.

5

Table 10-2

	Example 6	Comparative example 8
Treated hair:		
Gloss	9	1
Springness	8	2
Manageability	9	1

As shown in Table 10-2, the evaluated values of the hair treated by the hair spray of Example 6 are higher in every evaluation item of gloss, springness, and manageability of hair in comparison with the hair treated by the hair spray of Comparative Example 8. Therefore, N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxylpropyl hydrolyzed

10

15

collagen-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition contained in a hair spray clearly exerts excellent effects imparting gloss and springness to hairs, and improving manageability of hairs.

20

Example 7 and Comparative Examples 9 and 10

Three kinds of shampoo compositions shown in Table 11 were prepared. After shampooing hair 10 times by using each shampoo composition respectively, the hair smoothness

was evaluated and the properties of gloss, combability and manageability of hair were also evaluated.

In Example 7,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl

5 hydrolyzed

sericin-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition produced in Production Example 1 was used for a shampoo. In Comparative Example 9, in place of a silylated peptide-silane compound copolymer

10 composition, a silylated hydrolyzed sericin having about 500 of the number average molecular weight in its peptide portion was used for a shampoo; in Comparative Example 10, neither a silylated peptide-silane compound copolymer composition nor a silylated peptide were used for a shampoo.

15

Table 11

	Example	Comparative example	
	7	9	10
Silylated peptide-silane compound copolymer composition (70%) of Production Example 1	1.0	0	0
Silylated hydrolyzed sericin (20%)	0	5.0	0
Potassium (palm oil fatty acid) hydrolyzed collagen (35%) (a trade name of Promois ECP, manufactured by SEIWA KASEI Co., Ltd.)	50.0	50.0	50.0
(Palm oil fatty acid) diethanolamide	3.5	3.5	3.5
Paraoxybenzoic acid ester-phenoxyethanol mixture (a trade name of Seisept H, manufactured by SEIWA KASEI Co., Ltd.)	0.5	0.5	0.5
Perfume	Appropriate amount	Appropriate amount	Appropriate amount
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part	Volume adjustable totally 100 part

The hairs were treated for evaluation by following steps:

- 5 Preparing three bundles of hair, each bundle being the length of 15 cm and the weight of 1.5 g;
Washing a bundle with 2% aqueous solution of sodium polyoxyethylene(3) lauryl ether sulfate, and then rinsing it with streaming tap water; and then
- 10 Drying it with a hair dryer to provide a hair bundle for

shampoo treatment;

Getting wet said hair bundle for shampoo treatment with hot water;

Washing each hair bundle with 1 g of each of shampoos prepared
5 in Example 7, Comparative Example 9 and 10 respectively;
Rinsing it with streaming hot water; and then drying it with a hair dryer;

Repeating above procedures, which are shampooing, rinsing and drying with hair dryer, in 10 cycles; and then

10 Each of hair bundles was served to the hair smoothness evaluation test.

The hair smoothness was evaluated by measuring the friction force of the treated hair surface with the friction feeling tester (a trade name of KES-SE, made by Katotech
15 Co., Ltd.) under the conditions of humidity of $40 \pm 1\%$ and temperature $22 \pm 1^\circ\text{C}$.

The smoothness (, or roughness) is represented in said tester with the value of average deviation of friction coefficient sensed by a friction sensor traversing the
20 specified distance on the sample surface, wherein the unit of said value is dimensionless and the value indicates "more smoothness" when the value becomes smaller. In the present test, measurement was carried out 10 times per one sample and the average of obtained values was defined as the
25 measurement value.

Then, by using the same bundles of hair, the properties of gloss, combability and manageability of hair were sensory evaluated by 10 sensory-panelists in rating the properties

as follows: best (point 2), second best (point 1) and worse (point 0); and then the average of rated points was defined as the evaluation value. The results of evaluations are shown in Table 12 in terms of averaged value.

5

Table 12

	Example	Comparative example	
	7	9	10
Hair smoothness (The average deviation of friction coefficient)	0.0040	0.0045	0.0051
Treated hair:			
Gloss	1.6	1.4	0
Combability	2.0	1.0	0
Manageability	1.8	1.2	0

As shown in Table 12, the average deviation value of friction coefficient, which indicates hair smoothness, of the hair shampooed by the shampoo of Example 7 containing the silylated peptide-silane compound copolymer composition is the smallest; and said value is 88.9% of that of the hair shampooed by the shampoo of Comparative Example 9 blended with the silylated peptide, and 78.4% of that of the hair shampooed by the shampoo of Comparative Example 10 containing neither the silylated peptide-silane compound copolymer composition nor the silylated peptide. According this result, N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed

20

sericin-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition, which is blended in Example 7,
clearly exerts effects to be sorbed on and makes smooth
the hair surface when said copolymer composition is
5 contained in a shampoo.

Moreover, according to the sensory evaluation, the
evaluated values of the hair shampooed by the shampoo of
Example 7 are higher in every evaluation item such as gloss,
combability and manageability of hair in comparison with
10 the hairs shampooed by the shampoos of Comparative Example
9 and 10. Therefore, the silylated peptide-silane compound
copolymer composition blended in a shampoo of Example 7
clearly exerts excellent effects imparting gloss to hairs,
improving combability and manageability of hairs after
15 being shampooed.

Example 8 and Comparative Example 11

Two kinds of shampoo compositions shown in Table 13
were prepared. After shampooing hairs with each shampoo
20 composition respectively, the gloss, smoothness,
combability and manageability of hair for shampooed hair
were sensory evaluated.

In Example 8,
N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
25 hydrolyzed
collagen-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition of the silylated peptide-silane
compound copolymer composition produced in Production

Example 2 was used. In Comparative Example 11, in place of a silylated peptide-silane compound copolymer composition, a silylated hydrolyzed collagen having about 500 of the number average molecular weight in its peptide
5 portion was used.

Table 13

	Example	Comparative example
	8	11
Silylated hydrolyzed peptide-silane compound copolymer composition (70%) of Production Example 2	0.2	0
Silylated hydrolyzed collagen (20%)	0	0.7
Monosodium N-lauroyl-L-glutamate	15.0	15.0
Coconut fatty acid diethanolamide	3.5	3.5
Stearyltrimethyl ammonium chloride (30%)	3.3	3.3
Paraoxybenzoic acid ester-phenoxyethanol mixture (a trade name of Seisept H, manufactured by SEIWA KASEI Co., Ltd.)	0.5	0.5
Perfume	Appropriate amount	Appropriate amount
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part

The sensory evaluation test for the above described shampoo was carried by the half-head method, in which 10 sensory-panelists (consisting of 7 females and 3 males) combed their hair in two sides, that is, right and left sides, and then shampooed one of sides with the shampoo of Example 8 and the rest side with the shampoo of Comparative Example 11. Although the amount of shampoo used was different depending on the volume and length of their hair, they

shampooed once a day for seven days, then evaluated the properties of gloss, smoothness, combability and manageability of hair about that which of shampoo of Example 8 and Comparative Example 11 is better or both are not different (, or almost same).

The result is shown in Table 14 in the numbers of the panelists who answered as Example 8 being better, who answered as Comparative Example 11 being better and who answered as there being no difference.

Table 14

	Numbers answered as Example 8 better	Numbers answered as Comparative example 11 better	Numbers answered as no difference
For shampooed hair			
Gloss	9	0	1
Smoothness	10	0	0
Combability	10	0	0
Manageability	7	0	3

As shown in Table 14, the number of the panelists who answered as Example 8 being better is largest in every evaluation item, particularly, on the properties of smoothness and combability, every panelist answered as Example 8 being better. According this result,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxylpropyl hydrolyzed

collagen-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition is clearly excellent in the effects

to provide smoothness and combability to hair, moreover clearly exerts effects of imparting gloss to hairs and imparting manageability of hairs after hair being shampooed.

5

Example 9 and Comparative Example 12 and 13

Three kinds of shampoo compositions shown in Table 15 were prepared. After shampooing hairs in 10 cycles with each shampoo composition respectively, the hair smoothness was evaluated and the properties of gloss, combability and manageability of hair were also evaluated.

In Example 9,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed

15 silk-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition of the silylated peptide-silane compound copolymer composition produced in Production Example 3 was used for a shampoo. In Comparative Example 12, in place of a silylated peptide-silane compound
20 copolymer composition, a silylated hydrolyzed silk having about 600 of the number average molecular weight in its peptide portion was used for a shampoo; in Comparative Example 13, neither a silylated peptide-silane compound copolymer composition nor a silylated peptide were used
25 for a shampoo.

Table 15

	Example	Comparative example	
	9	12	13
Silylated peptide-silane compound copolymer composition (70%) of Production Example 3	0.1	0	0
Silylated hydrolyzed silk (20%)	0	0.4	0
Cocoylamidopropyl-dimethyl glycine (30%)	17.1	17.1	17.1
Polyoxyethylene(2) dodecyleter sulfate	8.3	8.3	8.3
Polyoxyethylene(100EO) sorbitanmonolaurate	7.5	7.5	7.5
Paraoxybenzoic acid ester-phenoxyethanol mixture (a trade name of Seisept H, manufactured by SEIWA KASEI Co., Ltd.)	0.5	0.5	0.5
Perfume	Appropriate amount	Appropriate amount	Appropriate amount
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part	Volume adjustable totally 100 part

By using shampoos obtained by above Example 9, Comparative Example 12 and 13 respectively, each bundle of hair having the length of 15 cm and the weight of 1.5 g was washed with 0.5 g of shampoo and hot water, and then was rinsed with hot water, and then was dried with a hair dryer. After repeating this procedure in 10 cycles, with the same manner carried in Example 7, the smoothness of hair surface was measured with the friction feeling tester, and then the properties of gloss, combability and

manageability of hair were sensory evaluated by 10 female sensory-panelists under same rating rule applied in Example 7. The results of those evaluations are shown in Table 16 in average values.

5

Table 16

	Example	Comparative example	
	9	12	13
Hair smoothness (The average deviation of friction coefficient)	0.0043	0.0046	0.0050
Treated hair:			
Gloss	1.7	1.3	0
Combability	1.9	1.1	0
Manageability	1.8	1.2	0

As shown in Table 16, the average deviation value of friction coefficient, which indicates hair smoothness, of the hair shampooed by the shampoo of Example 9 containing the silylated peptide-silane compound copolymer composition is the smallest; and said value is 93.5% of that of the hair shampooed by the shampoo of Comparative Example 12 containing the silylated peptide, and 82.7% of that of the hair shampooed by the shampoo of Comparative Example 13 containing neither the silylated peptide-silane compound copolymer composition nor the silylated peptide. According this result,

N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed silk-dimethyldiethoxysilane-octyltriethoxysilane

copolymer composition, which is blended in Example 9, clearly exerts effects to be sorbed on and makes smooth the hair surface when said copolymer composition is contained in a shampoo.

5 Moreover, according to the sensory evaluation, the evaluated values of the hair shampooed by the shampoo of Example 9 are higher in every evaluation item such as gloss, combability and manageability of hair in comparison with the hairs shampooed by the shampoos of Comparative Example 10 12 and 13. Therefore, the silylated peptide-silane compound copolymer composition blended in a shampoo of Example 9 clearly exerts excellent effects imparting gloss to hairs, improving combability of hairs and imparting manageability of hairs after being shampooed.

15

Example 10 and Comparative Example 14 and 15

First compositions of hair coloring agents (oxidation type) having composition as shown in Table 17 were prepared, and evaluated about dyeability, and gloss, springness, 20 combability and shampoo fastness of the dyed hairs.

In Example 10,
N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
hydrolyzed
sericin-dimethyldiethoxysilane-octyltriethoxysilane
25 copolymer composition produced in Production Example 1 was used. In Comparative Example 14, in place of a silylated peptide-silane compound copolymer composition, aminomethylaminopropylsiloxane/dimethylsiloxane

copolymer emulsion (SM8702C (trade name), manufactured by Dow Corning Toray Silicone Co.,Ltd.), an amino modified silicone, was used; in Comparative Example 15 (Control, blank), neither a silylated peptide-silane compound
5 copolymer composition nor a silicone were used.

Table 17

	Example	Comparative example	
	10	14	15
Silylated peptide-silane compound copolymer composition (70%) of Production Example 1	3.0	0	0
Amino modified silicone (40%)	0	5.3	0
p-phenylenediamine	0.8	0.8	0.8
p-aminophenol	0.2	0.2	0.2
o-aminophenol	1.0	1.0	1.0
Resolcinol	1.6	1.6	1.6
Isopropanol	9.0	9.0	9.0
Propyleneglycol	5.5	5.5	5.5
Oleic acid	15.0	15.0	15.0
Aqueous ammonia (28%)	10.0	10.0	10.0
Sodium sulfite	0.1	0.1	0.1
Oleyl alcohol	10.0	10.0	10.0
Edetate disodium	0.4	0.4	0.4
Bis-2-hydroxyethyl-sorbitaneamine	9.4	9.4	9.4
Hydroxyethyl stearylamine	6.0	6.0	6.0
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part	Volume adjustable totally 100 part

A second composition having composition as shown in Table 17-2 was prepared for the first composition of Example 10 and Comparative examples 14 and 15.

Table 17-2

Stearic acid	1.0
Glycerin monostearate	1.5
Polyoxyethylene(5) oleyl ether	1.0
Hydrogen peroxide(35%)	15.0
Sterilized ion-exchange water	Volume adjustable totally 100 part

The hairs were treated for evaluation by following
5 steps:

Preparing three bundles of hair, each bundle being the length
of 10 cm and the weight of 1 g;

Washing a bundle with 2% aqueous solution of
polyoxyethylene(3)laurylater sodium sulfate, and then
10 rinsing it with water;

Dipping it for 5 minutes in 10 g (for each bundle) of solution
prepared by mixing 6% aqueous hydrogenperoxide solution
and 2% aqueous ammonia in a weight ratio of 1:1 as bleaching
in order to make the comparison of dyeing power of dyed
15 hairs easier;

rinsing it with water;

Drying the hair; and then

Being subjected dyeing.

Dyeing was carried out by mixing same amounts of the
20 first composition and the second composition, coating 2
g of the mixture on each hair bundle, then leaving the bundle
for 15 minutes, rinsing it with hot water, and washing it

with 2% aqueous solution of sodium polyoxyethylene(3) laurylether sulfate. After dyeing was completed, the hair bundle was dried with a hair dryer, and the obtained hair bundle was evaluated about uniform dyeability, gloss, springness and combability by ten panelists.

In the evaluation, Example 10 and Comparative example 14 were compared with Comparative example 15 (Control), and evaluated according to the following rating.

10 Rating

+3: Much better than Control

+2: Better than Control

+1: A little better than Control

0: no difference was found with Control

15 -1: A little worse than Control

-2: Worse than Control

-3: Much worse than Control

A bundle of dyed hair with the hair coloring agent of Example 10, Comparative example 14 or Comparative example 15 was washed with 2% aqueous solution of sodium polyoxyethylene(3) laurylether sulfate, rinsed with water and dried with a hair dryer. After repeating above procedure 20 times, the washed hair bundle was evaluated about depth of color after washing, gloss, springness and combability by ten panelists according to the same rating as for the above evaluation. The results indicate shampoo fastness and are shown in Table 18 in terms of averaged

value of ten panelists.

Table 18

	Example	Comparative example
	10	14
Dyed hair before washing:		
Uniform dyeability	+2.3	+1.6
Gloss	+2.4	+1.5
Springness	+2.3	+1.0
Combability	+2.2	+0.6
Dyed hair after washing:		
Color Depth	+2.9	+0.8
Gloss	+1.9	+1.1
Springness	+2.3	+0.6
Combability	+1.9	+0.8

5 As shown in Table 18, the evaluated values of the hair dyed by the hair coloring agent of Example 10 are rated plus(+), that is, better than the hair dyed by the hair coloring agent of Comparative example 15 (Control), and are higher in every evaluation item in comparison with the
10 hair dyed by the hair coloring agent of Comparative example 14. Therefore,
N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed
sericin-dimethyldiethoxysilane-octyltriethoxysilane
15 copolymer composition, blended with the hair coloring agent, clearly exerts excellent effects giving uniform dyeability, suppressing damage of hair on dyeing (good combability indicates this fact), and imparting gloss, springness and combability to hairs. On the other hand, the hair dyed
20 by the hair coloring agent of Comparative example 14,

containing an amino modified silicone in place of a silylated peptide-silane compound copolymer composition, does not show high evaluated values, that is, improvements from Control are small, in every evaluation item such as gloss, springness and combability of hair. From this result, it is contemplated that the silicone blended in the hair coloring agent, is hard to be sorbed by hair or the silicone is easily washed away by washing after dyeing and the sorption amount is little, and, as the result, the effect is not realized.

Even after washing 20 times with shampoo, the evaluated values of the hair dyed by the hair coloring agent of Example 10 still higher in every evaluation items such as color depth, gloss, springness and combability of hair in comparison with the hair dyed by the hair coloring agent of Comparative example 14. From this result, it is clear that silylated peptide-silane compound copolymer composition blended in the hair coloring agent of Example 10 improve shampoo fastness of dyed hair.

Example 11 and Comparative Example 16 and 17

First compositions of hair coloring agents having composition as shown in Table 19 were prepared, and evaluated about uniform dyeability, and gloss, springness and combability of the dyed hairs.

In Example 11,
N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxylpropyl hydrolyzed

collagen-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition produced in Production Example 2 was
used. In Comparative Example 16, in place of a silylated
peptide-silane compound copolymer composition,
5 dimethylpolysiloxane copolymer emulsion (SH200C-100cs
(trade name), manufactured by Dow Corning Toray Silicone
Co., Ltd.), an amino modified silicone, was used; in
Comparative Example 17 (Control, blank), neither a
silylated peptide-silane compound copolymer composition
10 nor a silicone were used.

Table 19

	Example	Comparative example	
	11	16	17
Silylated peptide-silane compound copolymer composition (70%) of Production Example 2	5.0	0	0
Dimethylpolysiloxane	0	3.5	0
Nitro-p-phenylenediamine	0.4	0.4	0.4
4-amino-2-nitrophenol	0.4	0.4	0.4
o-aminophenol	0.1	0.1	0.1
Resorcinol	0.1	0.1	0.1
Polyvinylmethyle(7)nonyl phenyl ether	10.0	10.0	10.0
Oleic acid	5.0	5.0	5.0
Oleyl alcohol	10.0	10.0	10.0
Propyleneglycol	10.0	10.0	10.0
Isopropanol	10.0	10.0	10.0
Ascorbic acid	0.5	0.5	0.5
Aqueous ammonia (28%)	To adjust pH 9.5	To adjust pH 9.5	To adjust pH 9.5
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part	Volume adjustable totally 100 part

Second compositions of hair coloring agents having composition as shown in Table 19-2 were prepared.

Table 19-2

	Example	Comparative example	
	11	16	17
Hydrogen peroxide (35%)	15.0	15.0	15.0
Dimethylpolysiloxane	0.1	0.1	0.1
Nitro-p-phenylenediamine	84.9	84.9	84.9

Three bundles of hair, each bundle being the length of 10 cm and the weight of 1 g, were prepared, and bleached according to the same manner as in Example 10. Then dyeing was carried out using the hair coloring agent of Example 11, Comparative example 16 or Comparative example 17.

Dyeing was carried out by mixing same amounts of the first composition and the second composition, coating 2 g of the mixture on each hair bundle, then leaving the bundle for 20 minutes, rinsing it with hot water, and washing it with 2% aqueous solution of sodium polyoxyethylene(3) laurylether sulfate. After dyeing was completed, the hair bundle was dried with a hair dryer, and the obtained hair bundle was evaluated about uniform dyeability, gloss, springness and combability by ten panelists according to the same rating as in Example 10.

Further, a bundle of dyed hair with the hair coloring agent of Example 11, Comparative example 16 or Comparative example 17 was washed with 2% aqueous solution of sodium polyoxyethylene(3) laurylether sulfate, rinsed with water and dried with a hair dryer. After repeating above

procedure 20 times, the washed hair bundle was evaluated about depth of color after washing, gloss, springness and combability by ten panelists according to the same rating as in Example 10. The results indicate shampoo fastness and are shown in Table 20 in terms of averaged value of ten panelists.

Table 20

	Example	Comparative example
	11	16
Dyed hair before washing:		
Uniform dyeability	+2.3	+1.3
Gloss	+2.4	+1.1
Springness	+2.0	+0.9
Combability	+2.8	+0.5
Dyed hair after washing:		
Color Depth	+2.8	+0.6
Gloss	+2.0	+0.9
Springness	+2.1	+0.5
Combability	+2.0	+1.0

As shown in Table 20, the evaluated values of the hair dyed by the hair coloring agent of Example 11 blended with N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed collagen-dimethyldiethoxysilane-octyltriethoxysilane copolymer composition are rated plus(+), that is, better than the hair dyed by the hair coloring agent of Comparative example 17 (Control), and are higher in every evaluation item in comparison with the hair dyed by the hair coloring agent of Comparative example 16 blended with an amino modified silicone. Therefore, the silylated peptide-silane

compound copolymer composition blended in the hair coloring agent of Example 11 clearly exerts excellent effects giving uniform dyeability, suppressing damage of hair on dyeing, and imparting gloss, springness and combability to hairs.

5 Even after washing 20 times with shampoo, the evaluated values of the hair dyed by the hair coloring agent of Example 11 still higher in every evaluation items in comparison with the hair dyed by the hair coloring agent of Comparative example 16. From this result, it is clear that silylated
10 peptide-silane compound copolymer composition blended in the hair coloring agent of Example 11 improve shampoo fastness of dyed hair.

Example 12 and Comparative Example 18 and 19

15 First compositions of hair coloring agents having composition as shown in Table 21 were prepared, and evaluated about uniform dyeability, and gloss, springness and combability of the dyed hairs.

In Example 12,

20 N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl hydrolyzed

silk-dimethyldiethoxysilane-octyltriethoxysilane
copolymer composition produced in Production Example 3 was used. In Comparative Example 18, in place of a silylated
25 peptide-silane compound copolymer composition, octamethylcyclotetrasiloxane (SH244 (trade name), manufactured by Dow Corning Toray Silicone Co., Ltd.) was used; in Comparative Example 19 (Control, blank), neither

a silylated peptide-silane compound copolymer composition nor a silicone were used.

Table 21

	Example	Comparative example	
	12	18	19
Silylated peptide-silane compound copolymer composition (70%) of Production Example 3	1.5	0	0
Octamethylcyclotetrasiloxane	0	1.05	0
p-phenylenediamine	1.0	1.0	1.0
Propyleneglycol	10.0	10.0	10.0
Ammonium tioglycolate (50%)	0.5	0.5	0.5
Polyoxyethylene(6)stearyl ether	1.0	1.0	1.0
Edetate disodium	0.3	0.3	0.3
Aqueous ammonia (28%)	To adjust pH 10	To adjust pH 10	To adjust pH 10
Sterilized ion-exchange water	Volume adjustable totally 100 part	Volume adjustable totally 100 part	Volume adjustable totally 100 part

5

Second compositions of hair coloring agents having composition as shown in Table 21-2 were prepared.

Table 21-2

	Example	Comparative example	
	12	18	19
Hydrogen peroxide (35%)	6.0	6.0	6.0
Sterilized ion-exchange water	94.0	94.0	94.0

Three bundles of hair, each bundle being the length of 10 cm and the weight of 1 g, were prepared, and bleached according to the same manner as in Example 10. Then dyeing was carried out using the hair coloring agent of Example 12, Comparative example 18 or Comparative example 19.

Dyeing was carried out by mixing same amounts of the first composition and the second composition, coating 2 g of the mixture on each hair bundle, then leaving the bundle for 20 minutes, rinsing it with hot water, and washing it with 2% aqueous solution of sodium polyoxyethylene(3) lauryl ether sulfate. After dyeing was completed, the hair bundle was dried with a hair dryer, and the obtained hair bundle was evaluated about uniform dyeability, gloss, springness and combability by ten panelists according to the same rating as in Example 10.

Further, a bundle of dyed hair with the hair coloring agent of Example 12, Comparative example 18 or Comparative example 19 was washed with 2% aqueous solution of sodium polyoxyethylene(3) lauryl ether sulfate, rinsed with water and dried with a hair dryer. After repeating above procedure 20 times, the washed hair bundle was evaluated

about depth of color after washing, gloss, springness and combability by ten panelists according to the same rating as in Example 10. The results indicate shampoo fastness and are shown in Table 22 in terms of averaged value of
 5 ten panelists.

Table 22.

	Example	Comparative example
	12	18
Dyed hair before washing:		
Uniform dyeability	+2.5	+1.4
Gloss	+2.4	+1.4
Springness	+2.4	+0.9
Combability	+2.5	+1.1
Dyed hair after washing:		
Color Depth	+2.7	+0.6
Gloss	+2.2	+1.2
Springness	+2.6	+0.4
Combability	+2.2	+0.9

As shown in Table 22, the evaluated values of the hair
 10 dyed by the hair coloring agent of Example 12 blended with
 N-[2-hydroxy-3-(3'-methyldihydroxysilyl)propoxy]propyl
 hydrolyzed
 silk-dimethyldiethoxysilane-octyltriethoxysilane
 copolymer composition are rated plus(+), that is, better
 15 than the hair dyed by the hair coloring agent of Comparative
 example 19 (Control), and are higher in every evaluation
 item in comparison with the hair dyed by the hair coloring
 agent of Comparative example 18 blended with an amino
 modified silicone. Therefore, the silylated peptide-silane
 20 compound copolymer composition blended in the hair coloring

agent of Example 12 clearly exerts excellent effects giving uniform dyeability, suppressing damage of hair on dyeing, and imparting gloss, springness and combability to hairs.

Even after washing 20 times with shampoo, the evaluated
5 values of the hair dyed by the hair coloring agent of Example 12 still higher in every evaluation items in comparison with the hair dyed by the hair coloring agent of Comparative example 18. From this result, it is clear that silylated peptide-silane compound copolymer composition blended in
10 the hair coloring agent of Example 12 improve shampoo fastness of dyed hair.